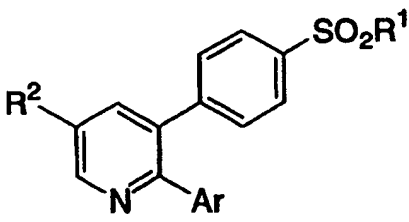




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07D 213/61, 213/50, C07B 43/00, A61K 31/44		A1	(11) International Publication Number: WO 98/47871
			(43) International Publication Date: 29 October 1998 (29.10.98)
(21) International Application Number: PCT/US98/08312 (22) International Filing Date: 14 April 1998 (14.04.98) (30) Priority Data: 60/045,642 18 April 1997 (18.04.97) US 9709686.1 13 May 1997 (13.05.97) GB (71) Applicant (for all designated States except US): MERCK & CO., INC. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): PYE, Philip, J. [GB/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). MALI-AKAL, Ashok [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). ROSSEN, Kai [DE/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). VOLANTE, Ralph, P. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). SAGER, Jess [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). (74) Common Representative: MERCK & CO., INC.; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).		(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, GW, HU, ID, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(54) Title: PROCESS FOR MAKING 2-ARYL-3-ARYL-5-HALO PYRIDINES USEFUL AS COX-2 INHIBITORS			
<div style="text-align: center;">  <p>(I)</p> </div>			
(57) Abstract The invention encompasses a process for making compounds of Formula (I) useful in the treatment of cyclooxygenase-2 mediated diseases.			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

TITLE OF THE INVENTION

PROCESS FOR MAKING 2-ARYL-3-ARYL-5-HALO PYRIDINES
USEFUL AS COX-2 INHIBITORS

5 BACKGROUND OF THE INVENTION

This invention concerns a process for making certain anti-inflammatory compounds. In particular, the application concerns a process for making compounds of formula I as disclosed hereinunder, which compounds are potent cyclooxygenase-2 inhibitors.

10 Non-steroidal, antiinflammatory drugs exert most of their antiinflammatory, analgesic and antipyretic activity and inhibit hormone-induced uterine contractions and certain types of cancer growth through inhibition of prostaglandin G/H synthase, also known as cyclooxygenase. Up until recently, only one form of cyclooxygenase had been characterized,
15 this corresponding to cyclooxygenase-1 or the constitutive enzyme, as originally identified in bovine seminal vesicles. Recently the gene for a second inducible form of cyclooxygenase (cyclooxygenase-2) has been cloned, sequenced and characterized from chicken, murine and human sources. This enzyme is distinct from the cyclooxygenase-1 which has now also been
20 cloned, sequenced and characterized from sheep, murine and human sources. The second form of cyclooxygenase, cyclooxygenase-2, is rapidly and readily inducible by a number of agents including mitogens, endotoxin, hormones, cytokines and growth factors. As prostaglandins have both physiological and pathological roles, we have concluded that the
25 constitutive enzyme, cyclooxygenase-1, is responsible, in large part, for endogenous basal release of prostaglandins and hence is important in their physiological functions such as the maintenance of gastrointestinal integrity and renal blood flow. In contrast, we have concluded that the inducible form, cyclooxygenase-2, is mainly responsible for the pathological
30 effects of prostaglandins where rapid induction of the enzyme would occur in response to such agents as inflammatory agents, hormones, growth factors, and cytokines. Thus, a selective inhibitor of cyclooxygenase-2 will have similar antiinflammatory, antipyretic and analgesic properties to a conventional non-steroidal antiinflammatory drug, and in addition would

inhibit hormone-induced uterine contractions and have potential anti-cancer effects, but will have a diminished ability to induce some of the mechanism-based side effects. In particular, such a compound should have a reduced potential for gastrointestinal toxicity, a reduced potential for renal side effects, a reduced effect on bleeding times and possibly a lessened ability to induce asthma attacks in aspirin-sensitive asthmatic subjects.

WO 96/24585 published August 15, 1996 and WO 96/10012, published April 4, 1996 disclose methods of making 2-aryl-3-aryl-pyridines. In the invention as descloses hereinunder, 2-aryl-3-aryl-pyridines are prepared in a simple to conduct, 1 step condensation from readily available starting materials. It is, therefore, surprisingly convenient and more efficient than the prevoiusly described procedure, in which the 2-aryl-3-aryl pyridine was constructed by serial stepwise addition of the aryl groups to the central pyridine ring. Moreover, the the process of the instant invention is also surprisingly superior in that expensive palladium reagents are not required nor is the combersome protection/de-protection sequense of the prior art process.

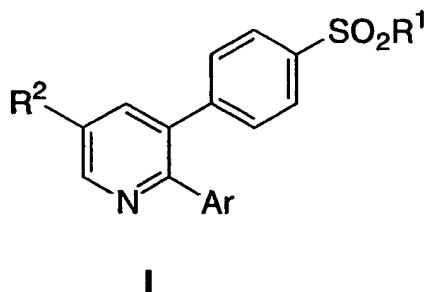
The preparation of 2-chloromalondialdehyde was first accomplished by Diekmann in 1904 (W. Dieckmann, L. Platz, *Ber. Deut. Chem. Ges.* 1904, 37, 4638). The chemistry of 2-halomalondialdehydes was thoroughly reviewed in 1975 (C. Reichardt and K. Halbritter, *Angew. Chem. Int. Ed.* 1975, 14, 86). This review does not mention a pyridine synthesis using these reagents. The only recorded use of 2-chloromalondialdehyde for the preparation of a pyridine is in a recent patent application (F.J. Urban, US 5,206,367 to Pfizer and Brackeen, M. and Howard, H.R. European Patent Application number 89307339.5 (EP 0 352 959) to Pfizer), where chloromalondialdehyde is first converted to 2,3-dichloroacrolein, which is subsequently condensed with the enamine derived from 1,3-cyclohexanedione to give the annulated pyridine in 28% yield.

A recent comprehensive review of pyridine synthesis and reactivity (D. Spitzner in *Methoden der Organischen Chemie* (Houben-Weyl), pages 286 to 686, Vol. E 7b, Editor R. P. Kreher, 1992, Georg

- Thieme Verlag) gives no examples for the use of halomalondialdehydes for the pyridine synthesis. Nitromalondialdehyde has been condensed with ethyl-2-amino-crotonate to give the 5-nitropyridine, albeit in lower yield (35-50%) (J.M. Hoffman et.al. *J. Org. Chem.* 1984, 49, 193 and P.E. Fanta, *J. Am. Chem. Soc.* 1953, 75, 737). The use of ethoxycarbonyl malondialdehyde derivatives leads to 5-ethoxycarbonyl pyridines (S. Torii et. al. *Synthesis*, 1986, 400).

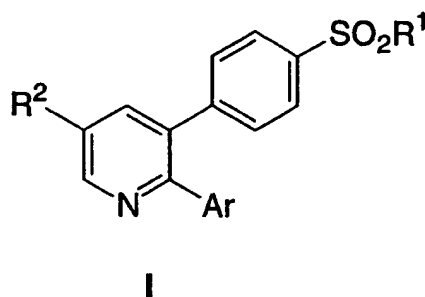
SUMMARY OF THE INVENTION

- 10 The invention encompasses a process for making compounds of Formula I useful in the treatment of inflammation and other cyclooxygenase-2 mediated diseases



15 DETAILED DESCRIPTION OF THE INVENTION

In a first aspect the invention encompasses a process for making compounds of Formula I useful in the treatment of inflammation and other cyclooxygenase-2 mediated diseases



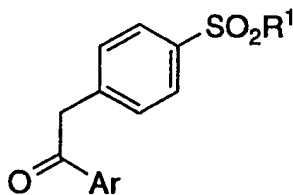
- 20 wherein:
R¹ is selected from the group consisting of

- (a) CH₃,
 (b) NH₂,
 (c) NHC(O)CF₃,
 (d) NHCH₃;
- 5 Ar is a mono-, di-, or trisubstituted phenyl or pyridinyl (or the N-oxide thereof), wherein the substituents are chosen from the group consisting of
- (a) hydrogen,
 (b) halo,
 (c) C₁₋₄alkoxy,
 10 (d) C₁₋₄alkylthio,
 (e) CN,
 (f) C₁₋₄alkyl,
 (g) C₁₋₄fluoroalkyl,
- R² is chosen from the group consisting of
- 15 (a) F, Cl, Br, I
 (b) CN,
 (c) azide,
- the process comprising:
 condensing a compound of formula A1



A1

under acidic conditions, and optionally in the presence of a non-reactive solvent and in the presence of an ammonium reagent, with compound A2



25

A2

to yield a compound of Formula I

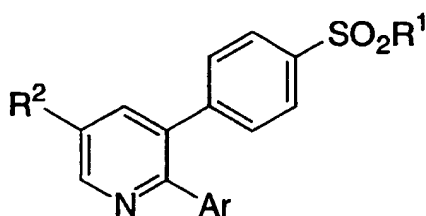
As will be appreciated by those of skill in the art, in the general case the reagents themselves provide the acidic condition. Therefore, the use of a non-reagent acid is not necessary. However, the addition of an acid, such as acetic or propionic or another carboxylic acid is within the scope of the invention.

For purposes of this specification non-reactive solvent includes tetrahydrofuran, dioxane, C₁-6alkanol, and toluene.

For purposes of this specification, the ammonium reagent is intended to include ammonia and ammonium salts such as ammonium acetate and ammonium propionate. Moreover a mixture ammonia reagent species is included in the term ammonia reagent.

The molar ratio of compound A1 to A2 can typically be varied from 2:1 to 1:2; preferably 1:1 to 1.5. Excess compound A1 is typically used. The molar ratio of compound A1 to ammonium reagent can typically be varied from 1:1 to 1:10. The reaction step may conveniently be conducted at a temperature range of 40 to 180°C; preferably 80 to 140°C and is allowed to proceed until substantially complete in from 2 to 18 hours; typically 6 to 12 hours.

In a second aspect the invention encompasses a process for making compounds of Formula I useful in the treatment of inflammation and other cyclooxygenase-2 mediated diseases



I

wherein:

- R¹ is selected from the group consisting of
- (a) CH₃,
 - (b) NH₂,
 - (c) NHC(O)CF₃,

(d) NHCH_3 ;

Ar is a mono-, di-, or trisubstituted phenyl or pyridinyl (or the N-oxide thereof), wherein the substituents are chosen from the group consisting of

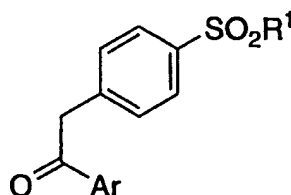
- 5 (a) hydrogen,
(b) halo,
(c) C_{1-4} alkoxy,
(d) C_{1-4} alkylthio,
(e) CN,
(f) C_{1-4} alkyl,
10 (g) C_{1-4} fluoroalkyl,

R^2 is chosen from the group consisting of

- (a) F, Cl, Br, I
(b) CN,
(c) azide,

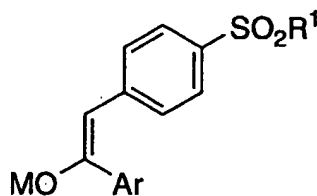
15 the process comprising:

(a) reacting a compound of formula A2



A2

20 in the presence of a second non-reactive solvent with a strong base to yield the enolate of formula B1



B1

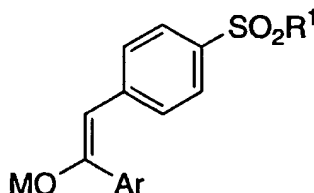
wherein M is potassium, lithium or sodium.

For purposes of this specification, the strong base shall include lithium, potassium or sodium diisopropylamide, lithium, potassium or sodium bis(trimethylsilyl)amide, lithium, potassium or sodium hydride, and lithium, potassium or sodium amide.

5 For purposes of this specification the second non-reactive solvent includes tetrahydrofuran, dioxane, toluene and ethers.

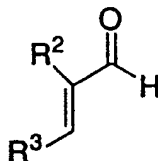
The molar ratio of compound A2 to base can typically be varied from 1:1 to 1:1.5. Excess base is typically used. The reaction step may conveniently be conducted at a temperature range of -80 to 40°C; preferably -10 to 20°C and is allowed to proceed until substantially complete in from 1 to 3 hours; typically 1 to 2 hours.

(b) reacting a compound of formula B1



B1

15 in the presence of a third non-reactive solvent with compound B2



B2

wherein R^3 is a leaving group such as tosyl, mesyl or halo.

20 which after heating in the presence of ammonia reagent, yields a compound of formula I.

For purposes of this reaction, the third non-reactive solvent shall include tetrahydrofuran, toluene and dioxane. The molar ratio of compound B1 to 2,3-dichloroacrolein can typically be varied from 1:1.5 to 1.5:1; preferably 1:1 to 1.5. Excess 2,3-dichloroacrolein is typically used.

25 The reaction step may conveniently be conducted at a temperature range of

0 to 80°C; preferably 20 to 50°C and is allowed to proceed until substantially complete in from 2 to 18 hours; typically 4 to 12 hours.

With regard to the both aspects of the invention, R² is preferably halogen, most preferably F or Cl, most preferably Cl. It is
5 preferable that R³ be the same as R².

With regard to both aspects of the invention a preferred sub-genus of formula I is that wherein Ar is a mono-, or disubstituted pyridinyl. Within this sub-genus, the 3-pyridinyl isomers are particularly preferred.

Again with regard to both aspects of the invention another
10 preferred sub-genus of formula I is that wherein R¹ is CH₃ or NH₂. Generally, CH₃ is preferred for COX-2 specificity and NH₂ is preferred for potency.

Again with regard to both aspects of the invention another preferred sub-genus of formula I is that wherein the Ar is unsubstituted or
15 substituted with CH₃.

The Compound of Formula I is useful for the relief of pain, fever and inflammation of a variety of conditions including rheumatic fever, symptoms associated with influenza or other viral infections, common cold, low back and neck pain, dysmenorrhea, headache, toothache,
20 sprains and strains, myositis, neuralgia, synovitis, arthritis, including rheumatoid arthritis degenerative joint diseases (osteoarthritis), gout and ankylosing spondylitis, bursitis, burns, injuries, following surgical and dental procedures. In addition, such a compound may inhibit cellular neoplastic transformations and metastatic tumor growth and hence can be
25 used in the treatment of cancer. Compounds of formula I may also be useful for the treatment of dementia including pre-senile and senile dementia, and in particular, dementia associated with Alzheimer Disease (ie Alzheimer's dementia).

By virtue of its high cyclooxygenase-2 (COX-2) activity and/or
30 its selectivity for cyclooxygenase-2 over cyclooxygenase-1 (COX-1) as defined above, compounds of formula I will prove useful as an alternative to conventional non-steroidal antiinflammatory drugs (NSAID'S) particularly where such non-steroidal antiinflammatory drugs may be contra-indicated such as in patients with peptic ulcers, gastritis, regional

enteritis, ulcerative colitis, diverticulitis or with a recurrent history of gastrointestinal lesions; GI bleeding, coagulation disorders including anemia such as hypoprothrombinemia, haemophilia or other bleeding problems (including those relating to reduced or impaired platelet
5 function); kidney disease (eg impaired renal function); those prior to surgery or taking anticoagulants; and those susceptible to NSAID induced asthma.

Compounds of the present invention are inhibitors of cyclooxygenase-2 and are thereby useful in the treatment of
10 cyclooxygenase-2 mediated diseases as enumerated above. This activity is illustrated by their ability to selectively inhibit cyclooxygenase-2 over cyclooxygenase-1. Accordingly, in one assay, the ability of the compounds of this invention to treat cyclooxygenase mediated diseases can be demonstrated by measuring the amount of prostaglandin E₂ (PGE₂)
15 synthesized in the presence of arachidonic acid, cyclooxygenase-1 or cyclooxygenase-2 and a compound of formula I. The IC₅₀ values represent the concentration of inhibitor required to return PGE₂ synthesis to 50 % of that obtained as compared to the uninhibited control. Illustrating this aspect, we have found that the Compounds of the Examples are more than
20 100 times more effective in inhibiting COX-2 than they are at inhibiting COX-1. In addition they all have a COX-2 IC₅₀ of 1 nM to 1 mM. By way of comparison, Ibuprofen has an IC₅₀ for COX-2 of 1 mM, and Indomethacin has an IC₅₀ for COX-2 of approximately 100 nM.

For the treatment of any of these cyclooxygenase mediated
25 diseases, compounds of formula I may be administered orally, topically, parenterally, by inhalation spray or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicles. The term parenteral as used herein includes subcutaneous injections, intravenous, intramuscular, intrasternal
30 injection or infusion techniques. In addition to the treatment of warm-blooded animals such as mice, rats, horses, cattle sheep, dogs, cats, etc., the compound of the invention is effective in the treatment of humans.

The invention will now be illustrated by the following non-limiting examples in which, unless stated otherwise:

- (i) all operations were carried out at room or ambient temperature, that is, at a temperature in the range 18-25°C; evaporation of solvent was carried out using a rotary evaporator under reduced pressure (600-4000 pascals: 4.5-30 mm. Hg) with a bath temperature of up to 60°C; the course of reactions was followed by thin layer chromatography (TLC) or High Pressure Liquid Chromatography (HPLC) and reaction times are given for illustration only; melting points are uncorrected and 'd' indicates decomposition; the melting points given are those obtained for the materials prepared as described; polymorphism may result in isolation of materials with different melting points in some preparations; the structure and purity of all final products were assured by at least one of the following techniques: TLC, mass spectrometry, nuclear magnetic resonance (NMR) spectrometry or microanalytical data; yields are given for illustration only; when given, NMR data is in the form of delta (δ) values for major diagnostic protons, given in parts per million (ppm) relative to tetramethylsilane (TMS) as internal standard, determined at 300 MHz or 400 MHz using the indicated solvent; conventional abbreviations used for signal shape are: s. singlet; d. doublet; t. triplet; m. multiplet; br. broad; etc.: in addition "Ar" signifies an aromatic signal; chemical symbols have their usual meanings; the following abbreviations have also been used v (volume), w (weight), b.p. (boiling point), m.p. (melting point), L (liter(s)), mL (milliliters), g (gram(s)), mg (milligrams(s)), mol (moles), mmol (millimoles), eq (equivalent(s)).

The following abbreviations have the indicated meanings:

Alkyl Group Abbreviations

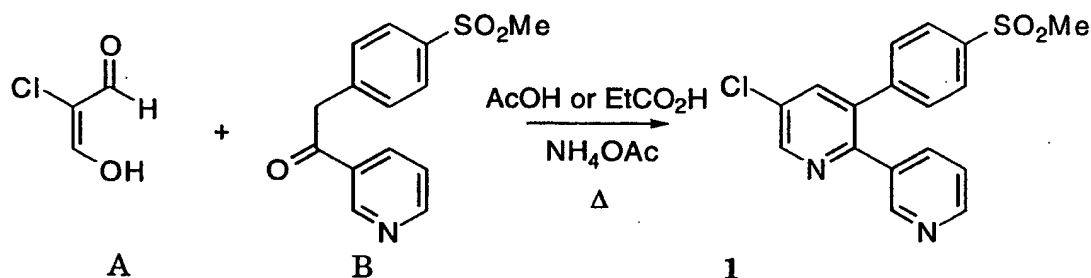
Me	=	methyl
Et	=	ethyl
n-Pr	=	normal propyl
i-Pr	=	isopropyl
n-Bu	=	normal butyl
i-Bu	=	isobutyl

s-Bu	=	secondary butyl
t-Bu	=	tertiary butyl
c-Pr	=	cyclopropyl
c-Bu	=	cyclobutyl
c-Pen	=	cyclopentyl
c-Hex	=	cyclohexyl

EXAMPLE 1

5-Chloro-3(methylsulfonyl)phenyl-2-(3-pyridyl)-pyridine; Compound 1

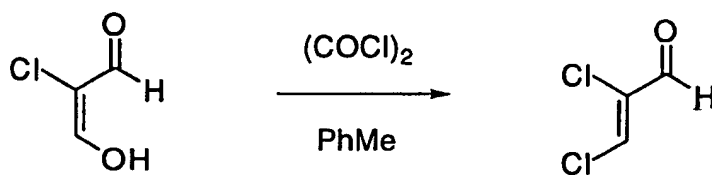
5



2-Chloromalondialdehyde	4.8 g (0.045 mol)
10 Ketone B	5.0 g (0.018 mol)
Propionic acid	30 mL
Ammonium Acetate	8.4 g (0.11 mol)

15 A mixture of ketone B (5.0 g), 2-chloromalondialdehyde (4.8 g) and ammonium acetate were heated to 130°C. The acetic acid produced was removed by distillation and heating continued at 136°C for 15 hours. The reaction mixture was basified with sodium carbonate, water was added and the product was extracted into dichloromethane (2 x 150 mL). The organic layers were carbon treated (Dowex), dried (MgSO₄) and the solvent removed to afford 1 as an off white solid (3.4 g, 55% yield).

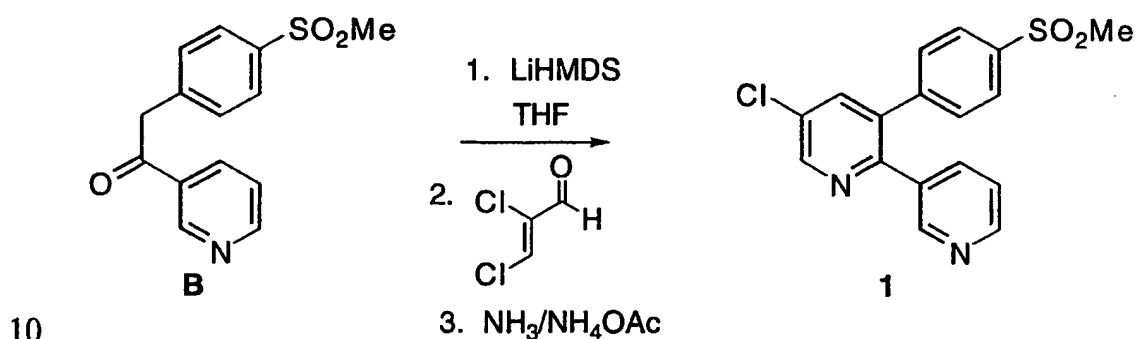
20



2-Chloromalondialdehyde	220 mg (2.1 mmol)
Oxalyl Chloride	180mL (2.1 mmol)
Toluene	3 mL
<i>N,N</i> -Dimethylformamide	20mL

5

N,N-dimethyl formamide was added to a slurry of 2-chloromalondialdehyde (220 mg) in toluene. Oxalyl chloride was added and the reaction mixture was stirred until complete dissolution occurred.



Ketone B	500 mg (1.8 mmol)
Lithium bis(trimethylsilyl)amide (1 M in THF)	1.8 mL (1.8 mmol)
Tetrahydrofuran	15 mL
2,3-Dichloroacrolein in toluene	2.1 mmol in 3 mL toluene
Ammonium acetate	1.0 g

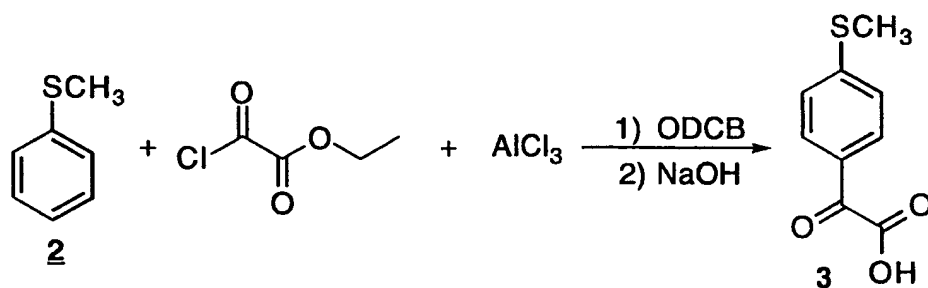
15 Lithium bis(trimethylsilyl)amide (1.8 mL; 1 M in THF) was added dropwise to ketone B (500 mg) in THF (15 mL) at -78°C. The reaction mixture was warmed to ambient temperature for 1 hour to form the lithium enolate of B (see the generic formula B1) before recooling to -78°C. A solution of 2,3-dichloroacrolein was added and the temperature allowed to warm to room temperature. After 1 hour ammonia gas was passed through the solution and after 30 minutes ammonium acetate (1 g) was added. The reaction mixture was warmed to 60°C for 1 hour and poured into aqueous sodium hydroxide (2 M; 100 mL). The product was extracted with dichloromethane (2 x 150 mL), dried (MgSO₄) and the solvent removed to afford 1 (500 mg; 80%).

30

PREPARATION OF STARTING MATERIALS

PREP 1SYNTHESIS OF 4-METHYLSULFONYLPHENYLACETIC ACID

5



	Thioanisole <u>2</u> (FW=124.2, d=1.058)	50.00g (0.403mol, 47.3mL)
	Ethyloxalyl chloride (FW=136.5, d=1.222)	82.43g (0.604mol, 67.5mL)
	Aluminum chloride (FW=133.3)	75.13g (0.564mol)
10	o-dichlorobenzene (ODCB)	112mL

The ethyloxalyl chloride and ODCB were charged to a flask equipped with an overhead mechanical stirrer and cooled to 0°C. The AlCl₃ was added slowly. The addition of the AlCl₃ was exothermic.

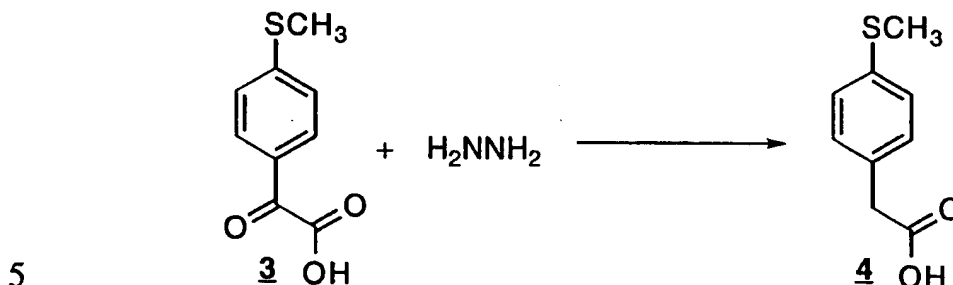
- 15 The thioanisole 2 was added dropwise via an addition funnel over 1.5 h. The reaction mixture rapidly turns a dark violet color. This addition was also exothermic.

- 20 After 1 h, the reaction was complete by HPLC. The reaction was quenched by the slow addition of 300mL of 1N HCl at 0°C. After warming to room temperature, water and ODCB (50mL each) were added. The layers were mixed and cut. The organic (bottom) phase was washed with 1 x 250mL water and then dried over MgSO₄.

- 25 This quench was also exothermic. The reaction mixture turned from dark violet to pale green during the quench. The dried ODCB solution was charged to a Morton flask equipped with mechanical stirring. A solution of 1N NaOH (800mL) was added. The biphasic mixture was stirred vigorously and heated to 50°C. Hydrolysis to 3 was complete in 2-3

h by HPLC. The product-containing aqueous phase was taken directly into the Wolf-Kishner reaction.

4-(Methylthio)phenylacetic acid



3 (in 1N NaOH solution)

(0.402mol)

Hydrazine (FW=32.1, 35wt% in water) 206.14g (2252mol, 204mL)

NaOH (5N solution)

5mL

10

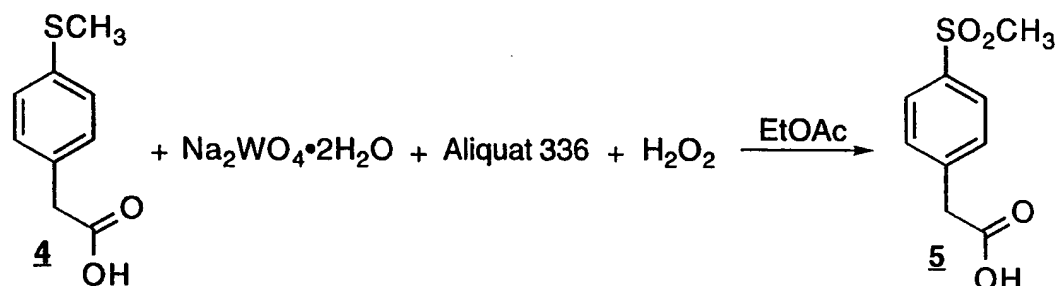
The hydrazine and NaOH were charged to a Morton flask equipped with mechanical stirring. After heating the hydrazine solution to 75°C, the solution of **3** in NaOH was added over 35-40 min. At the end of the addition the reaction mixture was brought to reflux for 5 days. HPLC showed the reaction to be ca. 95% complete at this point. The starting material was largely consumed in under 24 h, but a third peak which took several days to convert to **4**. The reaction was acidified with concentrated HCl to pH=1.5 and extracted with EtOAc (1 x 750mL and 1 x 250mL). The combined product-containing organic phases were washed 2 x 250mL 1N HCl.

15

20

On acidification, the reaction mixture turned bright yellow.

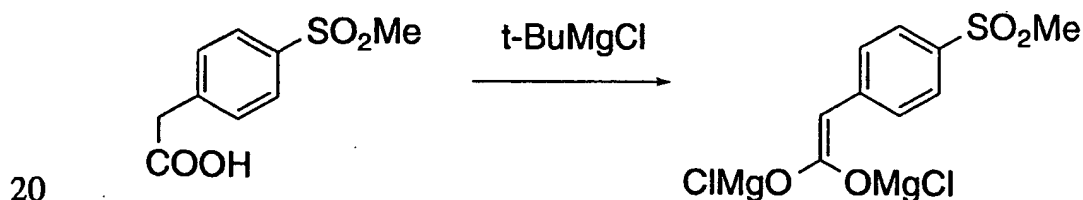
4-(Methanesulfonyl)phenylacetic acid

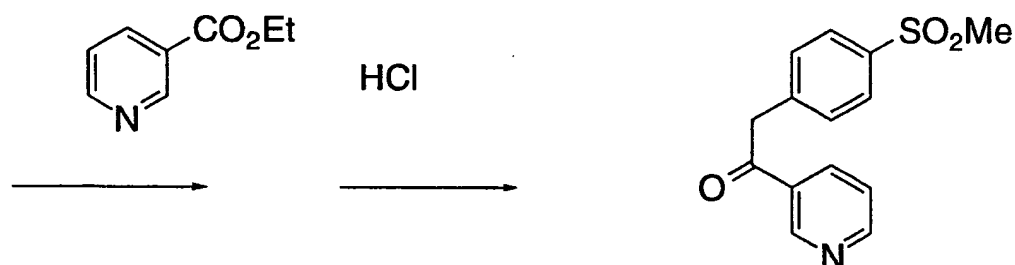


- 4-(Methylthio)phenylacetic acid 4 (FW=182.3) (0.402mol)
 Na₂WO₄·H₂O (FW=329.9) 2.64g (0.008mol)
 Aliquat 336 (FW=404) 8.08g (0.020mol)
 5 Hydrogen peroxide (FW=34.0, 30wt% in water) 136g (1.200mol, 123mL)

A flask equipped for mechanical stirring was charged with 3 (from reaction above, in EtOAc), Aliquat 336, and Na₂WO₄·2H₂O (dissolved in ca. 15mL H₂O). Hydrogen peroxide was added slowly via an addition funnel over ca. 30 min. Completion of reaction was checked by HPLC. The reaction was washed with 2 x 400mL H₂O and dried over MgSO₄. Quantification of product in the organic layer gave 61.29g 5 (71% yield from thioanisole). On concentration of the solution, a white solid precipitated. The slurry was filtered, and washed with hexanes. Recovery was 49.02g 5 (57% from thioanisole).

Ivanov-Claisen Condensation for the Preparation of
1-(3-pyridyl)-2-(4-methylsulfonylphenyl)-ethane-1-one





4-Methylsulfonylbenzyl-3-pyridylketone from ethyl nicotinate and 4-methylsulfonylphenylacetic acid

5

4-Methylsulfonylphenyl Acetic Acid (MW = 217)	10g (46.7 mmol)
<i>t</i> -Butyl magnesium chloride (1N/THF)	128.11ml (128.11mmol)
Ethyl nicotinate (MW = 151.2; d = 1.107)	5.54ml (39.4mmol)
THF	400ml

10

Phenyl acetic acid was dissolved in THF under nitrogen. 1.9 equivalents (88.73ml) of *t*-butyl magnesium chloride were added over 5 minutes to the solution. The Reaction was exothermic. The temperature rose from 20°C to 50°C. After addition of the first equivalent of *t*-butyl magnesium chloride, the solution turned red.

15

The reaction temperature was maintained at 50°C. After one hour, 0.5 equivalents of ethyl nicotinate were added. The solution turned yellow and a white precipitate formed. After one hour, 0.5 equivalents of *t*-butyl magnesium chloride were added at 50°C. The solution turned red. This sequence of addition was repeated using 0.25eq., 0.125eq., 0.0625eq. of ethyl nicotinate and *t*-butyl magnesium chloride. The reaction mixture was aged for 1 hour between each addition.

20

After the last addition, the reaction was quenched by adding the reaction mixture into vigorously stirred 2N hydrochloric acid (100ml). The solids at the bottom of the reaction mixture dissolved with effervescence when stirred in hydrochloric acid.

25

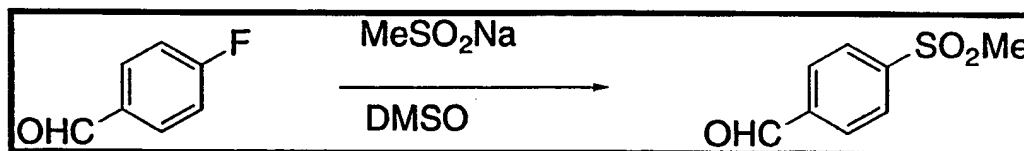
The pH of the aqueous phase of the reaction mixture was adjusted to 10 with sodium carbonate. LC assay showed 91% yield of ketone

30

Preparation of 4-Methylsulfonylbenzaldehyde

The preparation follows the procedure of Ulman JOC, pp4691 (1989).

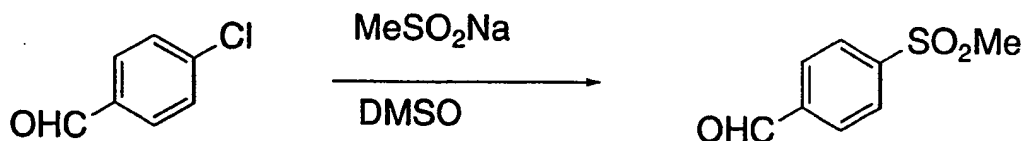
5 4-Methylsulfonylbenzaldehyde (2) from 4-Fluorobenzaldehyde



- 10 4-Fluorobenzaldehyde (MW = 124.11; d = 1.157) 23.3ml (217mmol)
Methanesulfinic acid, sodium salt (MW = 102.09) 24.23g (237mmol)
Methyl sulfoxide 170ml

Reagents were added to methyl sulfoxide and heated to 130°C for 18hrs. The sodium methanesulfinate was partially insoluble at RT but
15 went into solution at 130°C. Sodium fluoride precipitated out of solution. The reaction mixture was poured into 300ml water. The product precipitated out as a white solid. The reaction mixture was filtered. The product recovered was washed with 100ml water and 2x50ml methanol to remove methyl sulfoxide. The solvent was evaporated from the product
20 under reduced pressure affording 39.9g of 2 as a white powder (86% isolated yield). C^{13} -NMR (CDCl₃): 44.33, 128.25, 130.43, 139.70, 145.38, 190.72.

25 4-Methylsulfonylbenzaldehyde 2 from 4-Chlorobenzaldehyde

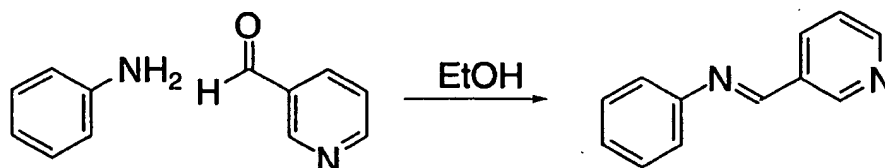


- 30 4-Chlorobenzaldehyde (MW = 140.57) 6.31g (45mmol)
Methanesulfinic acid, sodium salt (MW = 102.09) 7.5g (74mmol)
Methyl sulfoxide 50ml

Reagents were added to methyl sulfoxide and heated to 130°C for 18hrs.

The sodium methanesulfinate was partially insoluble at RT but went into solution at 130°C. Sodium chloride precipitated out of solution. The reaction mixture was poured into 100ml water. The product precipitated out as a white solid. The reaction mixture was filtered. The product recovered was washed with 50ml water and 2x25ml methanol to remove methyl sulfoxide. The solvent was evaporated from the product under reduced pressure affording 5.1g of 4-methylsulfonyl benzaldehyde as a white powder (62% isolated yield).

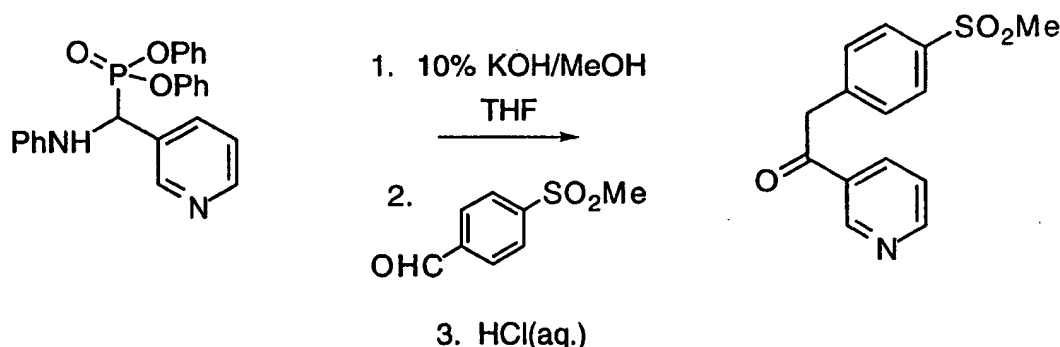
Horner/Wittig Route for the Preparation of 1-(3-pyridyl)-2-(4-methylsulfonylphenyl)-ethane-1-one



Ref: H. Zimmer, J. P. Bercz, *Liebigs Ann. Chem.* 1965, 686, 107-114.

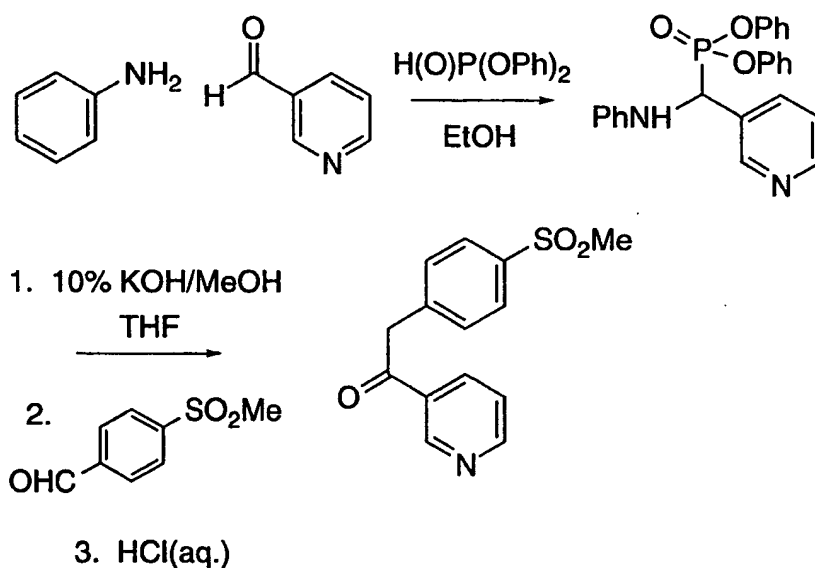
Aniline	89.4 g(0.96 mol)
3-pyridinecarboxaldehyde	102.8 g (0.96 mol)
Ethanol	150 mL
Diphenylphosphite	224.7 g (0.96 mol)

A solution of aniline in ethanol (50 mL) was added to a solution of 3-pyridine carboxaldehyde in ethanol (100 mL) at 0°C. After 2 hours diphenylphosphite was added and stirring was continued at room temperature for 18 hours. Methyltertbutylether (400 mL) was added to further precipitate the product which was filtered, washed (MTBE) and dried under vacuum to afford 320 g (80%) of the Pyridyl-amino diphenylphosphonate as a white solid. ¹³C NMR (CDCl₃):



	Pyridyl-amino diphenylphosphonate	14.0 g (0.034 mol)
	10% KOH in MeOH	23 mL (0.04 mol)
5	Tetrahydrofuran	150 mL
	4-methanesulfonylbenzaldehyde	5.6 g (0.03 mol)

10% KOH / MeOH (23 mL) was added over 10 minutes to a solution of phosphonate (14.0 g) in tetrahydrofuran at -45°C. After a further 10 minutes benzaldehyde was added in one portion and after 1 hour the reaction mixture was allowed to warm to ambient temperature. Aqueous hydrochloric acid (2N, 100 mL) was added and the solution was left standing for 18 hours. EtOAc (200 mL) and water (200 mL) were added and the organic layer discarded. The acid layer wash basified (pH = 9) with sodium carbonate and extracted with dichloromethane (2 x 150 mL). The organic layers were combined, dried (MgSO₄) and concentrated. Trituration with hexanes afforded 4-methylsulfonyl benzyl-3-pyridyl ketone as a pale yellow solid (6.3 g; 76%). ¹³C NMR (*D*-6 DMSO): 196.4, 153.6, 149.4, 140.8, 139.1, 135.7, 131.5, 130.9, 126.8, 123.9, 44.6 and 43.5 ppm.



	Aniline	4.47 g(0.05 mol)
5	3-pyridinecarboxaldehyde	5.36 g (0.05 mol)
	Methanol	10 mL
	Diphenylphosphite	11.2 g (0.05 mol)
	10% KOH in MeOH	28 mL (0.05 mol)
10	4-methanesulfonylbenzaldehyde	8.3 g (0.45 mol)

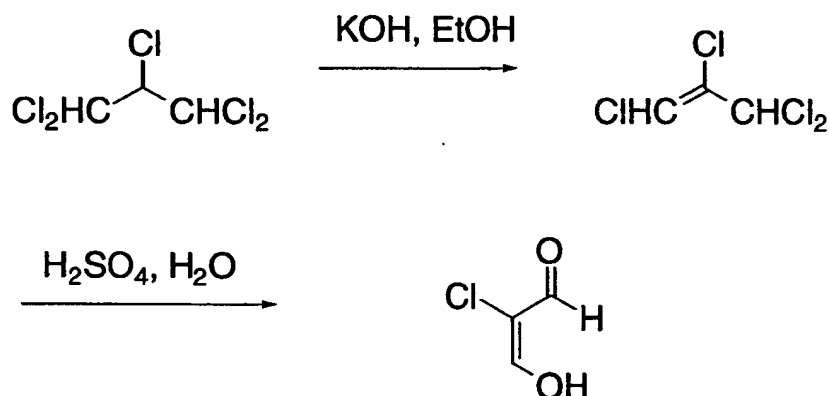
- A solution of aniline in methanol (5 mL) was added to a solution of 3-pyridine carboxaldehyde in methanol (5 mL) at 0°C. After 2 hours diphenylphosphite was added and stirring was continued at room temperature for 18 hours. THF (100 mL) was added and the reaction was cooled to -40°C. 10% KOH/methanol (28 mL) was added and after 30 minutes 4-methanesulfonylbenzaldehyde (8.3 g) was added. The reaction was allowed to warm to room temperature and stirred for 18 hours. EtOAc (200 mL) and water (200 mL) were added and the organic layer discarded.
- The acid layer was basified (pH = 9) with sodium carbonate and extracted with dichloromethane (2 x 150 mL). The organic layers were combined, dried (MgSO₄) and concentrated. Trituration with hexanes afforded 4-methylsulfonyl benzyl-3-pyridyl ketone as a pale yellow solid (9.7 g; 71%).

PREPARATION OF CHLOROMALONDIALDEHYDE

A number of routes are available for the preparation of chloromalondialdehyde.

5

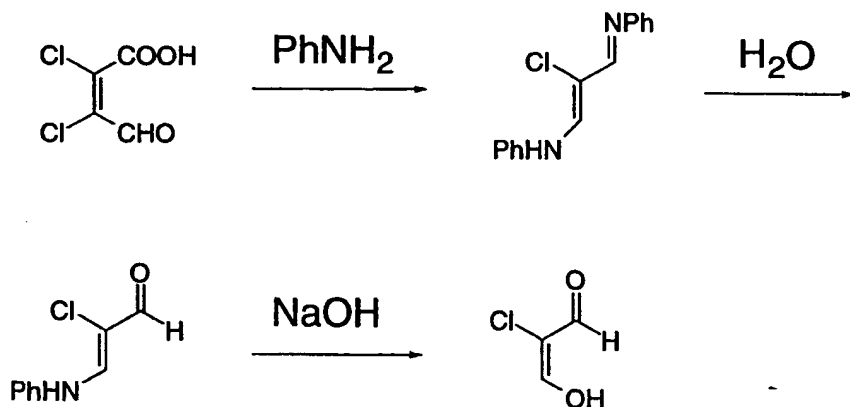
Preparation from 1,1,2,3,3-Pentachloropropane



10

A detailed experimental is published in Houben-Weyl-Muller: Methoden der Organischen Chemie, 4th Edit., Vol 7/1, Thieme Verlag, Stuttgart, 1954, page 119. The starting material 1,1,2,3,3-pentachloropropane is commercially available from Pfaltz and Bauer.

15 Preparation from Mucochloric Acid



The following is a slight variation of the original procedure of Dieckmann (*Ber. Deut. Chem. Ges.* 1904, 37, 4638).

5	Mucochloric acid	50.0 g (0.30 mol)
	Aniline	54 mL (0.60 mol)
	Water	1000 mL

To a solution of aniline in water at 85°C in a vigorously stirred 2 L flask was added mucochloric acid in small portions over 30 min.

- 10 On addition of the mucochloric acid, a yellow color develops, which quickly dissipated. The reaction mixture stayed heterogeneous and filtration of an aliquot after 30 min heating indicated completion of the reaction.

- 15 The reaction mixture was heated at 90°C for 60 min., cooled to 50°C and filtered. The filtercake was washed with 50 mL of 2N HCl and 100 mL of H₂O. The product was dried in a N₂ stream to give 57 g (100% yield) of 3-anilido-2-chloro-acrolein as a gray solid. ¹³C NMR (D₆-DMSO in ppm):108, 117, 124, 129, 140. 147, 182.

20	3-Anilido-2-chloro-acrolein	57 g (0.30 mol)
	5N NaOH solution	120 mL (0.6 mol)

A solution of 3-anilido-2-chloro-acrolein in 120 mL of 5N NaOH was heated to 100°C for 90 min. The dark black solution was extracted twice with 50 mL each of MTBE.

- 25 The first organic wash removed most of the dark color from the solution, and the second organic wash was only lightly colored.

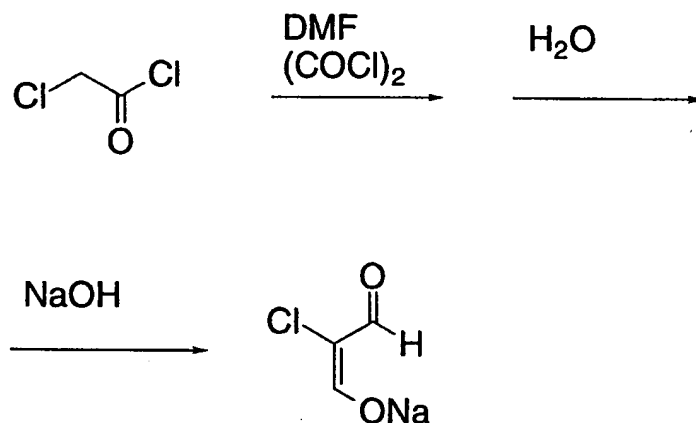
On cooling the aqueous phase, a crystalline precipitate formed. This product was the 3-chloromalondialdehyde Na salt.

- 30 The aqueous phase was acidified by the addition of 60 mL of 37% HCl solution. The aqueous phase was extracted (MTBE/THF 50/50, 400 mL total) and the combined organic phases were dried over MgSO₄. After treatment with Darco G60 and filtration through a plug of SiO₂, the solution was evaporated to give 19.6 g (62% overall yield) of 3-chloromalondialdehyde as a dark solid. Recrystallization from ca. 10 mL

of MTBE gave 11.13 g of pure chloromalondialdehyde as a tan solid. ^{13}C NMR ($\text{D}_6\text{-DMSO}$ in ppm): 113, 175 (broad).

Preparation from Chloroacetylchloride

5

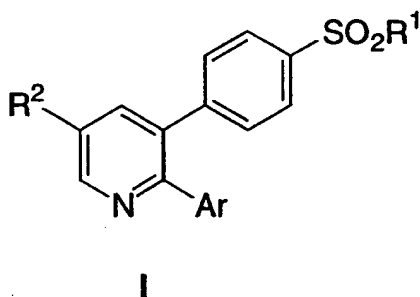


Arnold (*Collect. Czech. Chem. Commun.* 1961, 26, 3051) mentions the formation of 3-dimethylamino-2-chloro-acrolein by reaction of chloroacetic acid with the Vilsmeier reagent derived from POCl_3 and DMF. A variation and extension of his procedure prepares chloromalondialdehyde as its Na salt.

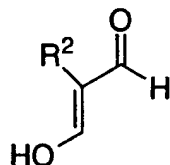
Oxalylchloride (280 mL, 3.2 mol) was added at 10°C to 1000 mL of DMF. The reaction was highly exothermic and a heavy precipitate formed. After a 2 h age, chloroacetylchloride (110 mL, 1.4 mol) was added and the reaction mixture was warmed to 75°C for 3 hours. Analysis of an aliquot by ^1H NMR indicated complete consumption of the chloroacetylchloride and the reaction mixture was quenched by addition into 1 L of H_2O . To the cooled solution was added 500 mL of a 50% NaOH solution. The reaction mixture is heated to reflux for 5 hours. On cooling a precipitate formed, which was filtered and washed with water. The tan solid was dried in a N_2 stream to give 84 g of a tan solid (54% yield).

WHAT IS CLAIMED

1. A process for making compounds of Formula I

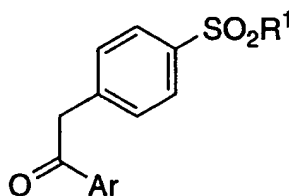


- 5 R^1 is selected from the group consisting of
 (a) CH_3 ,
 (b) NH_2 ,
 (c) $NHC(O)CF_3$,
 (d) $NHCH_3$;
- 10 Ar is a mono-, di-, or trisubstituted phenyl or pyridinyl (or the N-oxide thereof), wherein the substituents are chosen from the group consisting of
 (a) hydrogen,
 (b) halo,
 (c) C_{1-4} alkoxy,
15 (d) C_{1-4} alkylthio,
 (e) CN ,
 (f) C_{1-4} alkyl,
 (g) C_{1-4} fluoroalkyl,
- R^2 is chosen from the group consisting of
20 (a) F, Cl, Br, I
 (b) CN ,
 (c) azide,
- the process comprising:
 condensing a compound of formula A1



A1

under acidic conditions, and optionally in the presence of a non-reactive solvent and in the presence of an ammonium reagent, with compound A2



A2

to yield a compound of Formula I

2. A process according to Claim 1 wherein the non-reactive solvent is acetic acid.
3. A process according to Claim 1 wherein Ar is a mono- or di-trisubstituted 3-pyridinyl.
4. A process according to Claim 1 wherein R¹ is CH₃ or NH₂.
5. A process according to Claim 1 wherein Ar is a mono- or di-substituted 3-pyridinyl and the substituents are selected from the group consisting of
 - (a) hydrogen,
 - (b) halo,
 - (c) C₁-3alkoxy,
 - (d) C₁-3alkylthio,
 - (e) C₁-3alkyl,
 - (f) CF₃, and

(g) CN.

6. A process according to Claim 1 wherein R^1 is CH_3 or NH_2 ; and

5 Ar is a mono- or di-substituted 3-pyridinyl and the substituents are selected from the group consisting of

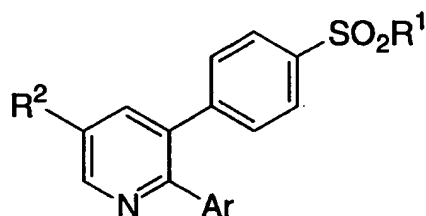
- (a) hydrogen,
(b) halo,
(c) C_{1-3} alkyl,
10 (d) CF_3 , and
(e) CN.

7. A process according to Claim 1 wherein R^2 is Cl;

15 R^1 is CH_3 or NH_2 ;

Ar is a mono-substituted 3-pyridinyl and the substituents are selected from the group consisting of hydrogen and C_{1-3} alkyl.

8. A process for making compounds of Formula I useful in
20 the treatment of inflammation and other cyclooxygenase-2 mediated diseases



I

wherein:

25 R^1 is selected from the group consisting of
(a) CH_3 ,
(b) NH_2 ,
(c) $NHC(O)CF_3$,

(d) NHCH_3 ;

Ar is a mono-, di-, or trisubstituted phenyl or pyridinyl (or the N-oxide thereof), wherein the substituents are chosen from the group consisting of

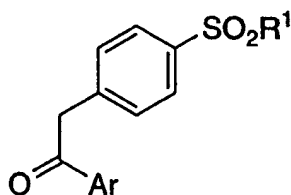
- 5 (a) hydrogen,
 (b) halo,
 (c) C_{1-4} alkoxy,
 (d) C_{1-4} alkylthio,
 (e) CN,
 (f) C_{1-4} alkyl,
 10 (g) C_{1-4} fluoroalkyl,

R^2 is chosen from the group consisting of

- (a) F, Cl, Br, I
 (b) CN,
 (c) azide,

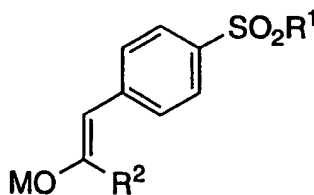
15 the process comprising:

(a) reacting a compound of formula A2



A2

20 in the presence of a second non-reactive solvent with a strong base to yield the enolate of formula B1

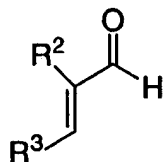


B1

wherein M is potassium, lithium or sodium, and

25 (b) reacting a compound of formula B1

in the presence of a third non-reactive solvent with compound B2



B2

- 5 wherein R³ is a leaving tosyl, mesyl or halo which after heating in the presence of ammonia reagent, yields a compound of formula I.
9. A process according to Claim 8 wherein Ar is a mono- or di-substituted 3-pyridinyl.
- 10 10. A process according to Claim 8 wherein R¹ is CH₃ or NH₂.
- 15 11. A process according to Claim 1 wherein Ar is a mono- or di-substituted 3-pyridinyl and the substituents are selected from the group consisting of
- 20 (a) hydrogen,
 (b) halo,
 (c) C₁₋₃alkoxy,
 (d) C₁₋₃alkylthio,
 (e) C₁₋₃alkyl,
 (f) CF₃, and
 (g) CN.
- 25 12. A process according to Claim 8 wherein R¹ is CH₃ or NH₂; and
- Ar is a mono- or di-substituted 3-pyridinyl and the substituents are selected from the group consisting of
- 30 (a) hydrogen,
 (b) halo,

- (c) C₁-3alkyl,
- (d) CF₃, and
- (e) CN.

5 13. A process according to Claim 8 wherein

R² is Cl;

R¹ is CH₃ or NH₂;

Ar is a mono-substituted 3-pyridinyl and the substituents are selected from the group consisting of hydrogen and C₁-3alkyl.

10

14. A process according to Claim 1 wherein R² is chloro.

15. A process according to Claim 8 wherein R³ is chloro.

15

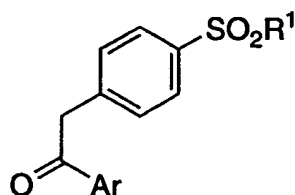
16. A process according to Claim 1 or 8 wherein the ammonium reagent is selected from ammonia and ammonium acetate.

17. A process according to Claim 8 wherein the strong base is lithium bis(trimethylsilyl)amide.

20

18. A process according to Claim 8 wherein the third non-reactive solvent is toluene.

19. A compound which is



25

R¹ is selected from the group consisting of

- (a) CH₃,
- (b) NH₂,
- (c) NHC(O)CF₃,
- (d) NHCH₃;

30

Ar is a mono-, di-, or trisubstituted phenyl or pyridinyl (or the N-oxide thereof), wherein the substituents are chosen from the group consisting of

- (a) hydrogen,
- (b) halo,
- 5 (c) C₁₋₄alkoxy,
- (d) C₁₋₄alkylthio,
- (e) CN,
- (f) C₁₋₄alkyl,
- (g) C₁₋₄fluoroalkyl,

10 R² is chosen from the group consisting of

- (a) F, Cl, Br, I
- (b) CN,
- (c) azide.

15 20. A compound according to Claim 19 wherein
R¹ is CH₃ or NH₂; and

Ar is a mono- or di-substituted 3-pyridinyl and the substituents are selected from the group consisting of

- 20 (a) hydrogen,
- (b) halo,
- (c) C₁₋₃alkyl,
- (d) CF₃, and
- (e) CN.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/08312

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D213/61 C07D213/50 C07B43/00 A61K31/44

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D C07B A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 22 21 546 A (CIBA-GEIGY AG) 16 November 1972 see example 13B ---	19,20
X,P	US 5 677 318 A (LAU CHEUK KUN) 14 October 1997 see example 1, step 2 ---	19
X	WO 96 38442 A (SEARLE & CO ; ROGERS KATHY L & HF (US); TALLEY JOHN J (US); SIKORSK) 5 December 1996 see example 9, step 2; example 14, step 1 ---	19
X	WO 96 25405 A (SEARLE & CO ; ROGERS KATHY L & LF (US); TALLEY JOHN J (US); BROWN D) 22 August 1996 see ex. 44, step 2; ex. 45, step 2; ex. 46, step 2 ---	19
-/--		

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

25 August 1998

Date of mailing of the international search report

01/09/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Bosma, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/08312

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 474 995 A (DUCHARME YVES ET AL) 12 December 1995 see example 6, step 1 ----	19
A	WO 96 24584 A (SEARLE & CO ;WEIER RICHARD M (US); LEE LEN F (US); PARTIS RICHARD) 15 August 1996 see scheme I -----	1-18
A,P	WO 98 03484 A (GAUTHIER JACQUES YVES ;MERCK FROSST CANADA INC (CA); DUBE DANIEL () 29 January 1998 see schemes 1 and 2 -----	1-18

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/08312

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE 2221546 A	16-11-1972	CH 579072 A	31-08-1976
		CH 561202 A	30-04-1975
		AR 194594 A	31-07-1973
		AR 198820 A	24-07-1974
		AR 198658 A	15-07-1974
		AR 195910 A	15-11-1973
		AT 319939 B	10-01-1975
		AT 319940 B	10-01-1975
		AT 319941 B	10-01-1975
		AT 316544 B	15-06-1974
		AU 472065 B	13-05-1976
		AU 4205572 A	15-11-1973
		BE 783244 A	10-11-1972
		CA 1012148 A	14-06-1977
		CH 561716 A	15-05-1975
		CH 561717 A	15-05-1975
		CH 561718 A	15-05-1975
		CS 181711 B	31-03-1978
		CS 181745 B	31-03-1978
		CS 181746 B	31-03-1978
		CS 181747 B	31-03-1978
		CY 936 A	23-06-1978
		DD 97654 A	12-05-1973
		FI 57407 B	30-04-1980
		FR 2137740 A	29-12-1972
		GB 1381031 A	22-01-1975
		HK 61477 A	16-12-1977
		KE 2796 A	06-01-1978
		NL 7206346 A	14-11-1972
		SE 405731 B	27-12-1978
		US 3940486 A	24-02-1976
		US 3929807 A	30-12-1975
		ZA 7202777 A	28-02-1973
		DE 2222638 A	23-11-1972
US 5677318 A	14-10-1997	NONE	
WO 9638442 A	05-12-1996	US 5643933 A	01-07-1997
		AU 6027996 A	18-12-1996
		EP 0828736 A	18-03-1998

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/08312

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9625405 A	22-08-1996	US 5633272 A	27-05-1997
		AU 4867196 A	04-09-1996
		BR 9607035 A	04-11-1997
		CA 2212836 A	22-08-1996
		CN 1181075 A	06-05-1998
		CZ 9702546 A	14-01-1998
		EP 0809636 A	03-12-1997
		FI 973292 A	10-10-1997
		NO 973711 A	06-10-1997
		PL 321814 A	22-12-1997
US 5474995 A	12-12-1995	AT 165825 T	15-05-1998
		AU 1269495 A	01-08-1995
		AU 1913297 A	14-08-1997
		AU 691119 B	07-05-1998
		AU 6197096 A	31-10-1996
		AU 6967494 A	17-01-1995
		BG 100247 A	28-06-1996
		BG 100350 A	31-12-1996
		BR 9406979 A	05-03-1996
		BR 9408478 A	26-08-1997
		CA 2163888 A	05-01-1995
		CA 2176973 A	25-12-1994
		CA 2176974 A	25-12-1994
		CA 2180651 A	13-07-1995
		WO 9500501 A	05-01-1995
		WO 9518799 A	13-07-1995
		CN 1125944 A	03-07-1996
		CN 1143365 A	19-02-1997
		CZ 9503146 A	15-05-1996
		DE 69410092 D	10-06-1998
		EP 0705254 A	10-04-1996
		EP 0739340 A	30-10-1996
		EP 0754687 A	22-01-1997
		EP 0822190 A	04-02-1998
		ES 2115237 T	16-06-1998
		FI 956119 A	19-12-1995
		FI 962800 A	06-09-1996
		HR 940373 A	31-12-1996
		HU 74070 A	28-10-1996

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/08312

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5474995 A		HU 74986 A	28-03-1997
		JP 9500372 T	14-01-1997
		JP 9506631 T	30-06-1997
		MX 9404749 A	31-01-1995
		NO 955256 A	23-02-1996
		NO 960393 A	09-07-1996
		PL 312196 A	01-04-1996
		SG 43841 A	14-11-1997
		SK 150295 A	08-01-1997
		US 5536752 A	16-07-1996
		US 5550142 A	27-08-1996
		US 5710140 A	20-01-1998
		ZA 9404501 A	13-03-1995
WO 9624584 A	15-08-1996	US 5686470 A	11-11-1997
		AU 4859396 A	27-08-1996
		EP 0808304 A	26-11-1997
WO 9803484 A	29-01-1998	AU 3331997 A	10-02-1998